

CLAIMS

We claim:

1. An isolated nucleic acid having a nucleotide sequence selected from the group consisting of:
  - (a) at least 10 consecutive nucleotides of SEQ ID NO: 1;
  - (b) at least 12 consecutive nucleotides of SEQ ID NO: 1;
  - (c) at least 14 consecutive nucleotides of SEQ ID NO: 1;
  - (d) at least 16 consecutive nucleotides of SEQ ID NO: 1;
  - (e) at least 18 consecutive nucleotides of SEQ ID NO: 1; and
  - (f) a sequence complementary to any one of the sequences of (a) –(e).
2. An isolated nucleic acid having a nucleotide sequence selected from the group consisting of:
  - (a) at least 10 consecutive nucleotides of SEQ ID NO: 3;
  - (b) at least 12 consecutive nucleotides of SEQ ID NO: 3;
  - (c) at least 14 consecutive nucleotides of SEQ ID NO: 3;
  - (d) at least 16 consecutive nucleotides of SEQ ID NO: 3;
  - (e) at least 18 consecutive nucleotides of SEQ ID NO: 3; and
  - (f) a sequence complementary to any one of the sequences of (a) –(e).
3. An isolated nucleic acid having a nucleotide sequence selected from the group consisting of:
  - (a) a sequence encoding a CatSper4 protein;
  - (b) a sequence encoding at least a transmembrane domain of a CatSper4 protein;
  - (c) a sequence encoding at least an extracellular loop of a CatSper4 protein;
  - (d) a sequence encoding at least a pore region of a CatSper4 protein;
  - (e) a sequence encoding at least an epitope of a CatSper4 protein having high predicted antigenicity; and
  - (f) a sequence complementary to any one of the sequences of (a)–(e).
4. An isolated nucleic acid as in claim 3 selected from the group consisting of:
  - (a) SEQ ID NO: 1;

- (b) SEQ ID NO: 3;
- (c) a sequence encoding a polypeptide comprising residues 46-75, 86-98, 108-137, 155-202, 209-233 and 242-272 of SEQ ID NO: 2;
- (d) a sequence encoding a polypeptide comprising residues 48-73, 81-111, 113-134, 156-204, 208-233 and 245-273 of SEQ ID NO: 4;
- (e) a sequence encoding a polypeptide comprising residues 76-85, 138-154 and 234-242 of SEQ ID NO: 2;
- (f) a sequence encoding a polypeptide comprising residues 74-80, 135-155 and 234-244 of SEQ ID NO 4;
- (g) a sequence encoding a polypeptide comprising residues 222-231 of SEQ ID NO: 2;
- (h) a sequence encoding a polypeptide comprising residues approximately residues 222-231 of SEQ ID NO: 4;
- (i) a sequence encoding a polypeptide comprising a high predicted antigenicity epitope of SEQ ID NO: 2;
- (j) a sequence encoding a polypeptide comprising a high predicted antigenicity epitope of SEQ ID NO: 4; and
- (k) a sequence complementary to any one of the sequences of (a)–(j).

5. An isolated nucleic acid encoding a polypeptide having at least 80% amino acid sequence identity with a polypeptide selected from the group consisting of:

- (a) a CatSper4 protein;
- (b) at least a transmembrane domain of a CatSper4 protein;
- (c) at least an extracellular loop of a CatSper4 protein; and
- (d) at least a pore region of a CatSper4 protein.

6. An isolated nucleic acid encoding a polypeptide having at least 80% amino acid sequence identity with a CatSper4 protein and having CatSper4 activity in a cell capable of expressing CatSper4 activity.

7. An isolated nucleic acid comprising a regulatory element having at least 80% nucleotide sequence identity to at least 100 consecutive nucleotides selected from SEQ ID NO: 5;

wherein said regulatory element is capable of promoting transcription of a coding sequence operably joined thereto in a mammalian cell in which a CatSper4 gene can be expressed.

8. An isolated nucleic acid comprising a nucleotide sequence that hybridizes to at least a portion of a nucleic acid of SEQ ID NO: 1 or SEQ ID NO: 3 under conditions including a wash step of 1.0 x SSC at 65°C.
9. An isolated nucleic acid as in claim 8 wherein said nucleic acid encodes a polypeptide having CatSper4 activity.
10. A nucleic acid comprising:
  - (i) a nucleotide sequence encoding a polypeptide having CatSper4 activity, wherein said nucleic acid hybridizes to at least a portion of a nucleic acid of SEQ ID NO: 1 or SEQ ID NO: 3 under conditions including a wash step of 1.0 x SSC at 65°C; and
  - (ii) a heterologous regulatory region operably joined to said sequence such that said sequence is expressed.
11. A nucleic acid comprising:
  - (i) a nucleotide sequence encoding a polypeptide having at least 80 percent amino acid sequence identity with an amino acid sequence of SEQ ID NO: 2 or 4; and
  - (ii) a heterologous regulatory region operably joined to said sequence such that said sequence is expressed.
12. A kit for detecting at least a portion of a CatSper4 nucleic acid comprising an isolated nucleic acid of any one of claims 1-7 and a means for detecting said isolated nucleic acid.
13. A kit as in claim 12 wherein said means for detecting said isolated nucleic acid comprises a detectable label bound thereto.

14. A kit as in claim 12 wherein said means for detecting said isolated nucleic acid comprises a labeled secondary nucleic acid which specifically hybridizes to said isolated nucleic acid.
15. A vector comprising an isolated nucleic acid of any one of claims 1-11.
16. A vector comprising a genetic construct capable of expressing a nucleic acid of any one of claims 3-11.
17. A vector as in claim 16 wherein said nucleic acid is operably joined to an exogenous regulatory region.
18. A vector as in claim 16 wherein said nucleic acid is operably joined to heterologous coding sequences to form a fusion vector.
19. A vector comprising an isolated nucleic acid of any one of claims 3-11.
20. A vector comprising an isolated nucleic acid of any one of claims 3-11 operably joined to a reporter gene.
21. A cell transformed with a nucleic acid of any one of claims 3-11.
22. A cell transformed with a genetic construct capable of expressing a nucleic acid of any one of claims 3-11.
23. A cell as in claim 22 wherein said nucleic acid is operably joined to heterologous coding sequences to encode a fusion protein.
24. A cell as in claim 22 wherein said cell is selected from the group consisting of bacterial cells, yeast cells, insect cells, nematode cells, amphibian cells, rodent cells, and human cells.

25. A cell as in claim 22 wherein said cell is selected from the group consisting of mammalian somatic cells, fetal cells, embryonic stem cells, zygotes, gametes, germ line cells and transgenic animal cells.

26. A non-human transgenic animal, wherein a genetic construct has introduced a modification into a genome of said animal, or an ancestor thereof, and wherein said modification is selected from the group consisting of insertion of a nucleic acid encoding at least a fragment of a CatSper4 protein, inactivation of an endogenous CatSper4 gene, and insertion by homologous recombination of a reporter gene operably joined to CatSper4 regulatory elements.

27. An animal as in claim 26 wherein said modification is insertion of a nucleic acid encoding a polypeptide selected from the group consisting of a CatSper4 protein, at least a transmembrane domain of a CatSper4 protein, at least an extracellular loop of a CatSper4 protein, at least a pore region of a CatSper4 protein, and at least an epitope of a CatSper4 protein having high predicted antigenicity.

28. An animal as in claim 26 wherein said animal is selected from the group consisting of rats, mice, hamsters, guinea pigs, rabbit, dogs, cats, goats, sheep, pigs, and non-human primates.

29. A substantially pure protein preparation comprising a polypeptide selected from the group consisting of:

- (a) a CatSper4 protein;
- (b) at least a transmembrane domain of a CatSper4 protein;
- (c) at least an extracellular loop of a CatSper4 protein;
- (d) at least a pore region of a CatSper4 protein; and
- (e) at least an epitope of a CatSper4 protein having high predicted

antigenicity.

30. A substantially pure protein preparation as in claim 29 wherein said polypeptide is selected from the group consisting of:

- (a) SEQ ID NO: 2;
- (b) SEQ ID NO: 4;

- (c) residues 46-75, 86-98, 108-137, 155-202, 209-233 and 242-272 of SEQ ID NO: 2;
- (d) residues 48-73, 81-111, 113-134, 156-204, 208-233 and 245-273 of SEQ ID NO: 4;
- (e) residues 76-85, 138-154 and 234-242 of SEQ ID NO: 2;
- (f) residues 74-80, 135-155 and 234-244 of SEQ ID NO: 4;
- (g) residues 222-231 of SEQ ID NO: 2;
- (h) residues 222-231 of SEQ ID NO: 4;
- (i) a high predicted antigenicity epitope of SEQ ID NO: 2; and
- (j) a high predicted antigenicity epitope of SEQ ID NO: 4.

31. A substantially pure protein preparation comprising a polypeptide having at least 80% amino acid sequence identity with a polypeptide selected from the group consisting of:

- (a) a CatSper4 protein;
- (b) at least a transmembrane domain of a CatSper4 protein;
- (c) at least an extracellular loop of a CatSper4 protein; and
- (d) at least a pore region of a CatSper4 protein.

32. A substantially pure protein preparation comprising a polypeptide having at least 80% amino acid sequence identity with a CatSper4 protein and having CatSper4 activity in a cell capable of expressing CatSper4 activity.

33. A substantially pure antibody preparation comprising an antibody raised against a CatSper4 epitope.

34. A substantially pure antibody preparation as in claim 33 wherein said epitope has high predicted antigenicity.

35. A substantially pure antibody preparation as in claim 33 wherein said epitope comprises an amino acid sequence within the an amino acid sequence selected from the group consisting of high predicted antigenicity epitopes of SEQ ID NO: 2, and high predicted antigenicity epitopes of SEQ ID NO: 4.

36. A substantially pure antibody preparation as in any one of claims 33-35 wherein said antibody is a monoclonal antibody.
37. A substantially pure antibody preparation as in any one of claims 33-35 wherein said antibody is an antibody fragment selected from the group consisting of an Fab fragment, an F(ab')<sub>2</sub> fragment, an Fv fragment, and a single-chain Fv fragment (scFv).
38. A kit for detecting at least an epitope of a CatSper4 protein comprising an anti-CatSper4 antibody of any one of claims 33-37 and a means for detecting said antibody.
39. A kit as in claim 38 wherein said means for detecting said anti-CatSper4 antibody comprises a detectable label bound thereto.
40. A kit as in claim 38 wherein said means for detecting said anti-CatSper4 antibody comprises a labeled secondary antibody which specifically binds to said anti-CatSper4 antibody.
41. A method of identifying a potential modulator of CatSper4 activity comprising:  
    contacting a candidate compound with a cell expressing a CatSper4 protein;  
    measuring an indicator of CatSper4 activity in said cell;  
    determining whether said candidate compound caused an increase or decrease in said indicator relative to a reference level; and  
    identifying said candidate compound as a potential modulator of CatSper4 activity if said increase or decrease is significant.
42. A method as in claim 41 wherein said indicator is an indicator of the level of mRNA encoding said CatSper4 protein.
43. A method as in claim 41 wherein said indicator is an indicator of the level of CatSper4 protein.

44. A method as in claim 41 wherein said indicator is an indicator of cation flux across a membrane of said cell.
45. A method as in claim 41 wherein said indicator is an indicator of whole cell or channel currents of said cell.
46. A method as in any one of claims 41-45 wherein said cell has been transformed with a genetic construct capable of expressing a CatSper4 protein.
47. A method as in claim 41 wherein said cell is a mature sperm cell and said indicator is a measure of sperm motility.
48. A method of identifying a potential modulator of CatSper4 activity comprising:  
    contacting under physiological conditions a candidate compound with CatSper4 moiety comprising at least a structural domain of a CatSper4 protein;  
    measuring binding, if any, between said candidate compound and said CatSper4 moiety;  
    identifying said candidate compound as a potential modulator of CatSper4 activity if said binding is significant.
49. A method as in claim 48 wherein said CatSper4 moiety is a polypeptide selected from the group consisting of:  
    (a) a CatSper4 protein;  
    (b) at least a transmembrane domain of a CatSper4 protein;  
    (c) at least an extracellular loop of a CatSper4 protein; and  
    (d) at least a pore region of a CatSper4 protein.
50. A method of decreasing the fertility of a male subject comprising:  
    administering to said male a compound which decreases CatSper4 activity.
51. A method of causing reversible infertility in a male subject comprising:  
    administering to said male a compound which decreases CatSper4 activity.



52. A method of contraception comprising:  
administering to a male subject a compound which decreases CatSper4  
activity.
53. A method of contraception comprising:  
administering to a female subject a compound which decreases CatSper4  
activity.
54. A method as in any one of claims 50-53 wherein said compound is in a  
formulation selected from the group consisting of an injection, a transdermal patch, a  
bioerodable implant, a lubricant, a moisturizer, a foam, a jelly, and a sponge.
55. A method of contraception as in claim 53 wherein:  
said female subject is a mammal and said compound is administered into at  
least one of the vagina, uterus and fallopian tubes of said female.
56. A method as in any one of claims 50-53 wherein said compound is selected  
from the group consisting of a nucleic acid which is antisense to at least a portion of a  
CatSper4 gene and an antibody to a CatSper4 protein.
57. A method as in claim 56 wherein said compound is an antibody fragment  
selected from the group consisting of an Fab fragment, an F(ab')<sub>2</sub> fragment, an Fv  
fragment, and an scFv fragment.
58. A method as in any one of claims 50-53 wherein said subject is a mammal.
59. A method as in claim 58 wherein said mammal is selected from the group  
consisting of humans, dogs, cats, cows, sheep, horses, mice, rats, raccoons, and  
gophers.
60. A method as in claim 58 wherein said subject is selected from the group  
consisting of a fish, an amphibian and an insect.

61. Use of a compound which decreases CatSper4 activity in the preparation of a medicament for decreasing the fertility of a male subject.
62. Use of a compound which decreases CatSper4 activity in the preparation of a medicament for causing reversible infertility in a male subject.
63. Use of a compound which decreases CatSper4 activity in the preparation of a contraceptive for administration to a male.
64. Use of a compound which decreases CatSper4 activity in the preparation of a contraceptive for administration to a female.
65. A use as in any one of claims 61-64 wherein said compound is in a formulation selected from the group consisting of an injection, a transdermal patch, a bioerodable implant, a lubricant, a moisturizer, a foam, a jelly, and a sponge.
66. A use as in claim 64 wherein:  
said female subject is a mammal and said compound is administered into at least one of the vagina, uterus and fallopian tubes of said female.
67. A use as in any one of claims 61-64 wherein said compound is selected from the group consisting of a nucleic acid which is antisense to at least a portion of a CatSper4 gene and an antibody to a CatSper4 protein.
68. A use as in claim 67 wherein said compound is an antibody fragment selected from the group consisting of an Fab fragment, an F(ab')<sub>2</sub> fragment, an Fv fragment, and an scFv fragment.
69. A use as in any one of claims 61-64 wherein said subject is a mammal.
70. A use as in claim 69 wherein said mammal is selected from the group consisting of humans, dogs, cats, cows, sheep, horses, mice, rats, raccoons, and gophers.

71. A use as in claim 69 wherein said subject is selected from the group consisting of a fish, an amphibian and an insect.
72. A contraceptive preparation comprising a compound which decreases CatSper4 activity.
73. A preparation as in claims 72 wherein said compound is selected from the group consisting of a nucleic acid which is antisense to at least a portion of a CatSper4 gene and an antibody to a CatSper4 protein.
74. A preparation as in claim 72 wherein said preparation is in a formulation selected from the group consisting of an injection, a transdermal patch, a bioerodable implant, a lubricant, a moisturizer, a foam, a jelly, and a sponge.
75. A method of diagnosing a CatSper4-related disorder in a mammal comprising determining the presence or absence of a mutation in a CatSper4 gene.
76. A method as in claim 75 wherein said method comprises:  
determining at least a portion of a CatSper4 gene sequence and comparing said determined sequence to a reference sequence;  
wherein the presence or absence of differences between said determined sequence and said reference sequence indicate the presence or absence of mutations in said CatSper4 gene.
77. A method of diagnosing a CatSper4-related disorder comprising determining the presence or absence of a mutation in a CatSper4 protein.
78. A method as in claim 77 wherein said method comprises:  
determining at least a portion of a CatSper4 protein sequence and comparing said determined sequence to a reference sequence;  
wherein the presence or absence of differences between said determined sequence and said reference sequence indicate the presence or absence of mutations in said CatSper4 gene.

79. A method as in claim 78 wherein said determination comprises contacting at least a fragment of said CatSper4 protein with an antibody known to bind to a CatSper4 protein in which a mutation is known to be present or absent and detecting binding between said antibody and said fragment of said CatSper4 protein.
80. A method of diagnosing a CatSper4-related disorder in a mammal comprising measuring an indicator of CatSper4 activity in said cell; and comparing said measured indicator to a reference level.
81. A method as in claim 80 wherein said indicator is an indicator of the level of mRNA encoding said CatSper4 protein.
82. A method as in claim 80 wherein said indicator is an indicator of the level of CatSper4 protein.
83. A method as in claim 80 wherein said indicator is an indicator of cation flux across a membrane of said cell.
84. A method as in claim 80 wherein said indicator is an indicator of whole cell or channel currents of said cell.
85. A method as in any one of claims 75-84 wherein said disorder is CatSper4-related infertility.
86. A method of genotyping a subject with respect to a CatSper4 gene comprising:  
determining at least a portion of a CatSper4 gene sequence and comparing said determined sequence to a reference sequence;  
wherein the presence or absence of differences between said determined sequence and said reference sequence indicate the presence or absence of a genotype corresponding to said reference sequence.
87. A method of genotyping a subject with respect to a CatSper4 gene comprising:  
determining at least a portion of a CatSper4 protein sequence and comparing said determined sequence to a reference sequence;

wherein the presence or absence of differences between said determined sequence and said reference sequence indicate the presence or absence of a genotype corresponding to said reference sequence.

88. A method as in claim 87 wherein said determination comprises contacting at least a fragment of said CatSper4 protein with an antibody known to bind to a CatSper4 protein comprising said reference sequence and detecting binding between said antibody and said fragment of said CatSper4 protein.

89. A method of *in vitro* fertilization by sperm having decreased CatSper4 activity comprising:

- removing a zona pellucida from at least one ovum;
- contacting said ovum with at least one of said sperm; and
- allowing said sperm to fertilize said ovum.

90. A method of *in vitro* fertilization by sperm having decreased motility comprising:

- removing a zona pellucida from at least one ovum;
- contacting said ovum with at least one of said sperm; and
- allowing said sperm to fertilize said ovum.

91. A method of *in vitro* fertilization by sperm having decreased ability to penetrate a zona pellucida comprising:

- removing a zona pellucida from at least one ovum;
- contacting said ovum with at least one of said sperm; and
- allowing said sperm to fertilize said ovum.

92. A method of treating a subject characterized by infertility due to decreased CatSper4 activity comprising:

- transforming sperm or sperm progenitors of said subject with a genetic construct capable of expressing a CatSper4 protein; and
- using transformed sperm of said subject to fertilize an ovum.

93. A method of treating a subject characterized by infertility due to decreased CatSper4 activity comprising:  
administering a CatSper4 protein to sperm or sperm progenitors of said subject, whereby said protein provides CatSper4 activity in said sperm or spermatids;  
and  
using sperm bearing said administered CatSper4 protein to fertilize an ovum.
94. A method of diagnosing an anti-CatSper4 antibody-mediated infertility caused by anti-CatSper4 antibodies present in a female urogenital tract comprising:  
obtaining a sample of antibodies present in a female urogenital tract;  
contacting said sample of antibodies with at least a fragment of a CatSper4 protein; and  
detecting binding between said sample of antibodies and said fragment of a CatSper4 protein.
95. A method as in claim 94 wherein said CatSper4 fragment comprises a CatSper4 epitopes having high predicted antigenicity.
96. A method as in claim 95 wherein said epitope is included within a sequence selected from the group consisting of high predicted antigenicity epitopes of SEQ ID NO: 2, and high predicted antigenicity epitopes of SEQ ID NO: 4.
97. A method of treating an anti-CatSper4 antibody-mediated infertility caused by anti-CatSper4 antibodies present in a female urogenital tract, comprising:  
administering into said urogenital tract an agent which specifically binds to said anti-CatSper4 antibodies in an amount effective to inhibit binding between said anti-CatSper4 antibodies and a CatSper4 protein present on sperm in said urogenital tract.
98. A method as in claim 97 wherein said agent comprises at least fragment of a CatSper4 protein including an epitope having high predicted antigenicity.

99. A method as in claim 98 wherein said epitope is included within a sequence selected from the group consisting of high predicted antigenicity epitopes of SEQ ID NO: 2, and high predicted antigenicity epitopes of SEQ ID NO: 4.

100. A method as in claim 98 wherein said agent comprises an anti-idiotypic antibody against said anti-CatSper4 antibodies.

101. A method of conducting a drug discovery business comprising:

- (a) identifying, by the assay of claim 41, one or more agents which antagonize CatSper4 activity;
- (b) determining if an agent identified in step (a), or an analog thereof, inhibits at least one of sperm motility or egg penetrance;
- (c) conducting therapeutic profiling of an agent identified as an inhibitor in step (b) for efficacy and toxicity in one or more animal models; and
- (d) formulating a pharmaceutical preparation including one or more agents identified in step (c) as having an acceptable therapeutic profile.

102. The method of claim 101, further including the step of establishing a system for distributing the pharmaceutical preparation for sale, and optionally including establishing a sales group for marketing the pharmaceutical preparation.

103. A method of conducting a drug discovery business comprising:

- (a) identifying, by the assay of claim 41, one or more agents which agonize CatSper4 activity;
- (b) determining if an agent identified in step (a), or an analog thereof, increases at least one of sperm motility or egg penetrance;
- (c) conducting therapeutic profiling of an agent identified as an agonist in step (b) for efficacy and toxicity in one or more animal models; and
- (d) formulating a pharmaceutical preparation including one or more agents identified in step (c) as having an acceptable therapeutic profile;

104. The method of claim 101, further including the step of establishing a system for distributing the pharmaceutical preparation for sale, and optionally including establishing a sales group for marketing the pharmaceutical preparation.

105. The method of claim 103, wherein step (a) comprises identifying one or more agents which agonize the activity of wild type CatSper4.

106. The method of claim 103, wherein step (a) comprises identifying one or more agents which agonize the activity of a CatSper4 protein containing one or more mutations.

107. A method of conducting a reproductive medicine business comprising:

- (a) examining a sperm sample from a male patient, wherein said patient is experiencing a fertility problem;
- (b) determining if said sperm are characterized by at least one of a decrease in motility or a decrease in egg penetrance;
- (c) performing *in vitro* analysis to determine the efficacy of a CatSper4 agonist in increasing at least one of sperm motility or egg penetrance;
- (d) establishing a treatment regimen comprising administering an amount of a CatSper4 agonist effective to increase at least one of sperm motility or egg penetrance in said male.

108. The method of claim 107, further including a step wherein said male is monitored by a physician to evaluate improvement in fertility.

109. The method of claim 107, further including a step of billing the patient or the patient's health care provider.

110. A method of conducting a contraceptive medicine business comprising:

- (a) providing a pharmaceutical preparation discovered in claim X01, wherein said preparation inhibits the activity of CatSper4;
- (b) providing instructions to physicians and health care providers for the administration of an amount of said pharmaceutical preparation effective to inhibit the activity of CatSper4, wherein said effective amount is sufficient to prevent pregnancy.



111. The method of claim 110, further including the step of establishing a system for distributing the pharmaceutical preparation for sale, and optionally including establishing a sales group for marketing the pharmaceutical preparation.